APPENDIX B

COMPARISON OF CLAIM 1 OF 08/236,402 WITH THE PROPOSED COUNT

Appln. No. 08/236,402

Claim 1 (amended)

Count

A reagent for preparing a scintigraphic imaging agent,

A peptide comprising

a specific binding compound having a molecular weight of less than 10,000 daltons,

localize at a target site

a biological-function domain which causes the peptide to

the compound being covalently linked to

and

a radiolabel complexing moiety

a metal ion-binding domain

having a formula selected from the group consisting of: $I. \\ R^{1}\text{-CO-(amino acid)}^{1}\text{-(amino acid)}^{2}\text{-}Z$

which comprises the sequence Gly-Gly-Z or Gly-Gly-Z wherein Z is selected from the group consisting of cysteine, homocysteine, isocysteine, penicillamine, 2-mercaptoethylamine, 3-mercaptopropylamine and D-stereoisomers thereof.

wherein (amino acid)¹ and (amino acid)² are each independently any primary α - or β -amino acid that does not contain a thiol group; Z is selected from the group consisting of cysteine, homocysteine, isocysteine, penicillamine, 2-mercaptoethylamine and 3-mercaptopropylamine; R^1 is lower (C^1 - C^4) alkyl or covalent linkage to the compound;

wherein when Z is cysteine, homocysteine, isocysteine or penicillamine, Z comprises a carbonyl group covalently linked to a hydroxyl group, a NR³R⁴ group wherein R³ and R⁴ are each independently H or lower (C¹-C⁴) alkyl, an amino acid, or a peptide comprising 2 to 10 amino acids,

nd

=

Y-(amino acid)²-(amino acid)¹-NHR²

wherein Y is selected from the group consisting of cysteine, homocysteine, isocysteine, penicillamine, 2-mercaptoacetate and 3-mercaptopropionate; (amino acid)¹ and (amino acid)² are each independently any primary α - or β -amino acid that does not contain a thiol group; R^2 is selected from the group of H, a lower (C¹-C⁴) alkyl, a covalent linkage to the compound;

wherein when Y is cysteine, homocysteine, isocysteine or penicillamine, Y comprises an amino group covalently linked to -H, an amino acid, or a peptide comprising 2 to 10 amino acids; and

wherein the moiety is linked to the compound through R¹, R², a sidechain group of (amino acid)¹, a sidechain group of (amino acid)², an amino group of cysteine, homocysteine, isocysteine, or a carboxyl group of cysteine, homocysteine, isocysteine or penicillamine.

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